## IN THE CLAIMS

This listing of claims replaces all prior versions, and listings, in this application.

Claims 1-20 (canceled)

21. (currently amended) An enzyme solution comprising (a) an isolated β1,3-N-acetyl-D-galactosamine transferase which transfers N-acetyl-D-galactosamine to N-acetyl-D-glucosamine with β-1,3 linkage and has an amino acid sequence encoded by a nucleotide sequence that can hybridize to the complement of (i) SEQ ID NO:1 from nucleotide 106 to nucleotide 1503 or (ii) SEQ ID NO:3 from nucleotide 103 to nucleotide 1512 under high stringency conditions of hybridizing in 2x SSC and 50% formamide at 40°C and washing in 0.2x SSC and 0.1% SDS at 68°C; sharing at least 90% sequence identity with (i) SEQ ID NO: 2 from amino acid 189 to amino acid 500 or (ii) SEQ ID NO: 4 from amino acid 35 to amino acid 504, (b) buffer with a pH of at least 5.50 to 5.78 in MES buffer, a pH of around 5.0 in sodium cacodylate buffer, or a pH of around 7.4 to 7.5 in HEPES buffer; which is not from 6.2 to 6.6, and (c) divalent metal ion.

Claims 22-24 (canceled)

25. (currently amended) The enzyme solution of claim 21, wherein the  $\beta$ 1,3-N-acetyl-D-galactosamine transferase is encoded by a nucleotide sequence that can hybridize to the complement of SEQ ID NO: 1 from nucleotide 106 to nucleotide 1503 under high stringency conditions of <u>hybridizing hybridizating</u> in 2× SSC and 50% formamide at 40°C and washing in 0.2× SSC and 0.1% SDS at 68°C.

Claim 26 (canceled)

27. (previously presented) The enzyme solution of claim 21, wherein the  $\beta$ 1,3-N-acetyl-D-galactosamine transferase is encoded by a nucleotide sequence that can hybridize to

the complement of SEQ ID NO: 3 from nucleotide 103 to nucleotide 1512 under high stringency conditions of hybridizing in 2× SSC and 50% formamide at 40°C and washing in 0.2× SSC and 0.1% SDS at 68°C.

28. (currently amended) The enzyme solution of claim 21, wherein the pH of the buffer is at least 5.50 to 5.78 in MES buffer from about 5.0 to about 7.5.

29. (currently amended) The enzyme solution of claim 21, wherein the pH of the buffer is around 5.0 in sodium cacodylate buffer (i) greater than 5.0 and less than 6.2 or (ii) greater than 6.6 and less than 7.5.

30. (previously presented) The enzyme solution of claim 21, wherein the divalent metal ion is Mn<sup>2+</sup>, Co<sup>2+</sup>, or Mg<sup>2+</sup>.

31. (withdrawn) A process of using an enzyme solution of claim 21, wherein the process comprises catalyzing transfer of an N-acetyl-D-galactosamine (GalNAc) residue of a donor substrate to an N-acetyl-D-glucosamine (GlcNAc) residue of an acceptor substrate, wherein linkage between GalNAc and GlcNAc residues is a  $\beta$ 1,3 glycosidic linkage, in the enzyme solution.

Claims 32-34 (canceled)

35. (withdrawn) The process according to claim 31, wherein the  $\beta$ 1,3-N-acetyl-D-galactosamine transferase is encoded by a nucleotide sequence that can hybridize to the complement of SEQ ID NO: 1 from nucleotide 106 to nucleotide 1503 under high stringency conditions of hybridization in 2× SSC and 50% formamide at 40-50°C and washing in 0.2× SSC and 0.1% SDS at 68°C.

Claim 36 (canceled)

- 37. (withdrawn) The process according to claim 31, wherein the  $\beta$ 1,3-N-acetyl-D-galactosamine transferase is encoded by a nucleotide sequence that can hybridize to the complement of SEQ ID NO: 3 from nucleotide 103 to nucleotide 1512 under high stringency conditions of hybridization in 2× SSC and 50% formamide at 40-50°C and washing in 0.2× SSC and 0.1% SDS at 68°C.
- 38. (withdrawn/currently amended) The process according to claim 31, wherein the pH of the buffer is at least 5.50 to 5.78 in MES buffer from about 5.0 to about 7.5.
- 39. (withdrawn/currently amended) The process according to claim 31, wherein the pH of the buffer is around 5.0 in sodium cacodylate buffer (i) greater than 5.0 and less than 6.2 or (ii) greater than 6.6 and less than 7.5.
- 40. (withdrawn) The process according to claim 31, wherein the divalent metal ion is Mn<sup>2+</sup>, Co<sup>2+</sup>, or Mg<sup>2+</sup>.
- 41. (new) The enzyme solution of claim 21, wherein the  $\beta$ 1,3-N-acetyl-D-galactosamine transferase is comprised of the amino acid sequence shown in SEQ ID NO: 2.
- 42. (new) The enzyme solution of claim 21, wherein the  $\beta$ 1,3-N-acetyl-D-galactosamine transferase is comprised of the amino acid sequence shown in SEQ ID NO: 4.
- 43. (new) The enzyme solution of claim 21, wherein the  $\beta$ 1,3-N-acetyl-D-galactosamine transferase is comprised of the amino acid sequence from amino acid 189 to amino acid 500 of SEQ ID NO: 2.

- 44. (new) The enzyme solution of claim 21, wherein the  $\beta$ 1,3-N-acetyl-D-galactosamine transferase is comprised of the amino acid sequence from amino acid 35 to amino acid 504 of SEQ ID NO: 4.
- 45. (new) The process according to claim 31, wherein the  $\beta$ 1,3-N-acetyl-D-galactosamine transferase is comprised of the amino acid sequence shown in SEQ ID NO: 2.
- 46. (new) The process according to claim 31, wherein the  $\beta$ 1,3-N-acetyl-D-galactosamine transferase is comprised of the amino acid sequence shown in SEQ ID NO: 4.
- 47. (new) The process according to claim 31, wherein the  $\beta$ 1,3-N-acetyl-D-galactosamine transferase is comprised of the amino acid sequence from amino acid 189 to amino acid 500 of SEQ ID NO: 2.
- 48. (new) The process according to claim 31, wherein the  $\beta$ 1,3-N-acetyl-D-galactosamine transferase is comprised of the amino acid sequence from amino acid 35 to amino acid 504 of SEQ ID NO: 4.